

SEQUENCE LISTING

<110> Stanton, Jr., Vincent P.

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HAVING UTILITY IN DETERMINING THE TREATMENT
OF DISEASE

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<211> 3951
<212> DNA
<213> Homo sapiens

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<222> 166, 3432, 3682, 3937
<223> n = t or c
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 <212> DNA
 <213> Homo sapiens

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 <222> 175, 1067
 <223> n = g or a

<221> misc_feature
 <222> 341
 <223> n = c or g

<221> misc_feature
 <222> 791, 1997, 2618, 2653
 <223> n = t or c

<221> misc_feature
 <222> 1337
 <223> n = c or a

<221> misc_feature
 <222> 2107
 <223> nucleotide in position 2107 is g, or absent

<221> misc_feature
 <222> 2583
 <223> n = t or g

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<210> 5
<211> 3772
<212> DNA
<213> Homo sapiens

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<222> 431, 441
<223> n = a or g

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<221> misc_feature
<222> 498
<223> n = c or t

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<221> misc_feature
<222> 579, 599
<223> n = g or c

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<212> DNA

<213> Homo sapiens

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<221> misc_feature

<222> 1066

<223> n = t or c

<221> misc_feature
 <222> 1136
 <223> n = a or g

<221> misc_feature
 <222> 1497
 <223> n = t or a

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<213> Homo sapiens

<223> n = c or t

 $\langle 223 \rangle \quad n = g \text{ or } a$

<223> nucleotide at position 1045 is c, or absent

<223> nucleotide at position 1046 is t, or absent

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 <222> 2636, 5287
 <223> n = c or g

<221> misc_feature
 <222> 3118
 <223> n = g or t

<221> misc_feature
 <222> 3257, 4053
 <223> n = a or c

<221> misc_feature
 <222> 5440
 <223> n = t or a

<400> 11

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<213> Homo sapiens

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<222> 802, 1900
<223> n = c or t

<221> misc_feature
<222> 1747
<223> n = t or g

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 <213> Homo sapiens

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 <223> n = a or g

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 <222> 266
 <223> n = g or t

<221> misc_feature
 <222> 527
 <223> n = c or g

<221> misc_feature
 <222> 1217, 1647, 2282
 <223> n = c or t

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<223> n = c or t

<221> misc_feature

<222> 1289

<223> n = c or a

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<222> 771

<223> n = g or c

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<222> 883

<223> n = g or a

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<221> misc_feature

<222> 1052

<223> n = a or c

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